PAS

PREMIER ANALYTICAL SCIENCES

REPORT NO.

PAS-2026-5430

ISSUED 2026-11-19

PAGE 1 OF 1

REVISION 0 — CURRENT

CLIENT

Patrick Settles

CHAIN

RECEIVED 2026-11-10

CONDITION — INTACT, SEALED

ANALYZED 2026-11-18

RELEASED 2026-11-19 · 4:32 PM PST

BASIS — PER VIAL

METHODS & LIMITS

a · Fluorometric — Fluorometric assay (peptide content) (SOP-13); LOQ 20 µg/mL

b · HPLC-UV / ESI-QTOF-MS — USP/NF 621 Chromatography (SOP-13); LOQ 1 µg/mL

ACCEPTANCE

Purity: 97 % or higher.

Content: 50 to 150 % of the label claim.

Identity: conforms to the reference standard.

VERIFY

https://premieranalytical.com/verify/PAS-2026-5430

CERTIFICATE OF ANALYSIS

INDEPENDENT ANALYTICAL LABORATORY · MISSION HILLS, CALIFORNIA

Retatrutide

CAS 2381089-83-2 · LABEL CLAIM 30 MG · CLIENT DESIGNATION "RT30" · 1 VIAL, NUMBERED · MATERIAL AS STATED BY CLIENT

0 OF 1 SAMPLES PASS Nothing on this certificate was reportable across 1 sample. Each sample below states why.

SAMPLE 01 · LAB NO. PAS-26-01710 · RECEIVED INTACT, SEALED · DESIGNATION "RT30" **FLAGGED**

Retatrutide identity: DOES NOT CONFORM.
Retatrutide quantity (content) not reportable: Unable to quantify — excessive impurities.
Retatrutide purity not reportable: Unable to quantify — excessive impurities.

PURITY	% OF LABEL	IDENTITY
—	—	**DOES NOT CONFORM**
Unable to quantify — excessive impurities[b]	Unable to quantify — excessive impurities[a]	HPLC-UV / ESI-QTOF-MS[b]

ALSO MEASURED — Retatrutide — Unable to quantify — excessive impurities[a]



PHOTO 1 — AS RECEIVED



PHOTO 2 — LAB NO. AFFIXED

Released by — Donald Trujillo, Principal Chemist

released electronically; recorded on the verification register

RELEASED 2026-11-19 · 4:32 PM PST

REVISION 0 · STATUS: CURRENT

PAS-2026-5430 · SHA-256 1EC5 91DA CB9F B82A 61AB … · VERIFY: https://premieranalytical.com/verify/PAS-2026-5430 · PAGE 1 OF 1

Results relate only to the specific samples tested. Reproduce only in full. This certificate reports only the analyses listed, and its results are intended for research use only — the material tested is not approved for human or veterinary use, or for therapeutic, diagnostic, or clinical applications. Results should be interpreted by qualified professionals within the scope of the intended research. Accuracy and reliability of results may be influenced by sample integrity, handling, and other experimental variables. An endotoxin screen is not a sterility claim. Premier Analytical Sciences does not endorse any company or product listed in this register. A record here states what was measured, nothing more. Premier Analytical Sciences conducts its testing in accordance with ISO/IEC 17025 requirements; it is not an ISO/IEC 17025 accredited laboratory.

/// END OF REPORT ///